

From: [Biscoe, Melanie](#)
To: [Britton, Cathryn](#); [Fletcher, Rachel](#)
Subject: RE: Seresto Team Meeting - Prep to bring in FDA
Date: Wednesday, September 8, 2021 6:34:00 PM

I think Rachel's edit didn't save- so I changed it. I am done commenting on it.

Melanie L. Biscoe

Senior Regulatory Advisor, RMIB5
Pesticide Re-evaluation Division
US EPA Office of Pesticide Programs

From: Britton, Cathryn <Britton.Cathryn@epa.gov>
Sent: Wednesday, September 8, 2021 4:21 PM
To: Fletcher, Rachel <fletcher.rachel@epa.gov>
Cc: Biscoe, Melanie <Biscoe.Melanie@epa.gov>
Subject: RE: Seresto Team Meeting - Prep to bring in FDA
Thanks!

From: Fletcher, Rachel <fletcher.rachel@epa.gov>
Sent: Wednesday, September 08, 2021 4:21 PM
To: Britton, Cathryn <Britton.Cathryn@epa.gov>
Cc: Biscoe, Melanie <Biscoe.Melanie@epa.gov>
Subject: RE: Seresto Team Meeting - Prep to bring in FDA
I'm comfortable, so I made the edit.

From: Britton, Cathryn <Britton.Cathryn@epa.gov>
Sent: Wednesday, September 8, 2021 4:14 PM
To: Fletcher, Rachel <fletcher.rachel@epa.gov>
Cc: Biscoe, Melanie <Biscoe.Melanie@epa.gov>
Subject: FW: Seresto Team Meeting - Prep to bring in FDA
Rachel – are you comfortable providing an update on the petition for the FDA meeting? If so, please replace your name with “PRD” in the agenda. Thanks!

From: Herrick, Jacquelyn <Herrick.Jacquelyn@epa.gov>
Sent: Wednesday, September 08, 2021 2:15 PM
To: Shuler, Jamey <Shuler.Jamey@epa.gov>; Britton, Cathryn <Britton.Cathryn@epa.gov>; Biscoe, Melanie <Biscoe.Melanie@epa.gov>; Fletcher, Rachel <fletcher.rachel@epa.gov>; Miller, David <Miller.DavidJ@epa.gov>; Niman, Aaron <niman.aaron@epa.gov>; Recore, Shanna <Recore.Shanna@epa.gov>; Rossmeisl, Colleen <Rossmeisl.Colleen@epa.gov>; Breeden-Alemi, Julie <Breeden-Alemi.Julie@epa.gov>; Backus, Byron <Backus.Byron@epa.gov>; Andrews, Elizabeth <Andrews.Elizabeth@epa.gov>; Saunders, Jennifer <Saunders.Jennifer@epa.gov>; Leifer, Kerry <Leifer.Kerry@epa.gov>; Hospital, Jocelyn <Hospital.Jocelyn@epa.gov>; Walls, Cassi <Walls.Cassi@epa.gov>
Cc: Spatz, Dana <Spatz.Dana@epa.gov>; Dunbar, Anwar <Dunbar.Anwar@epa.gov>; Reaves, Elissa <Reaves.Elissa@epa.gov>; Echeverria, Marietta <Echeverria.Marietta@epa.gov>; Aubee, Catherine <Aubee.Catherine@epa.gov>; Vogel, Dana <Vogel.Dana@epa.gov>; Matuszko, Jan <Matuszko.Jan@epa.gov>; Di Salvo, Paul <DiSalvo.Paul@epa.gov>; Colby, Deanna <colby.deanna@epa.gov>; Kiely, Timothy <Kiely.Timothy@epa.gov>
Subject: RE: Seresto Team Meeting - Prep to bring in FDA

Hi all,

Per our discussion at the Seresto team meeting today, attached is the draft agenda for next week's meeting with FDA. Please comment and revise as you see fit. My day got a little out of hand, apologies for not getting this out earlier. Please take a look before **12pm tomorrow** if possible so we can follow up and solicit comment from the FDA.

Many thanks,

Jackie

Jacquelyn Herrick, PM 3

Invertebrate-Vertebrate Branch 1

Registration Division

Office of Pesticide Programs

U.S. Environmental Protection Agency

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herrick.jacquelyn@epa.gov

-----Original Appointment-----

From: Shuler, Jamey <Shuler.Jamey@epa.gov>

Sent: Thursday, September 02, 2021 9:55 AM

To: Shuler, Jamey; Britton, Cathryn; Biscoe, Melanie; Fletcher, Rachel; Miller, David; Niman, Aaron; Recore, Shanna; Rossmeisl, Colleen; Breeden-Alemi, Julie; Backus, Byron; Andrews, Elizabeth; Saunders, Jennifer; Herrick, Jacquelyn; Leifer, Kerry; Hospital, Jocelyn; Walls, Cassi

Cc: Spatz, Dana; Dunbar, Anwar; Reaves, Elissa; Echeverria, Marietta; Aubee, Catherine; Vogel, Dana; Matuszko, Jan; Di Salvo, Paul; Colby, Deanna; Kiely, Timothy

Subject: Seresto Team Meeting - Prep to bring in FDA

When: Wednesday, September 08, 2021 10:00 AM-11:00 AM (UTC-05:00) Eastern Time (US & Canada).

Where:

Hi Team,

FDA has agreed to meet in the next few weeks (09/16/2021), which grants us a little more time to prepare. Being that we are awaiting on decisions surrounding who will lead the science reviews, I have rescheduled today's meeting to next week **(09/08/2021)**. Have a great rest of the week!

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This discussion is regarding how we will loop in FDA and utilize them in our review (review specific studies? incidents? review our approach?). We were given word, that we will have 3 FDA reviewers to help with the science review. Additionally, we should solve who will review what and begin brainstorming how to use them. Thank you!

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